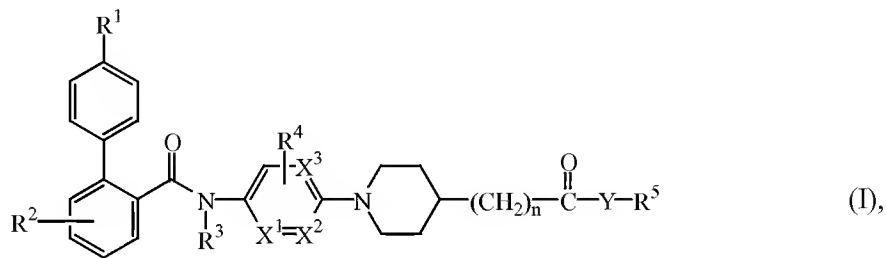


Listing of Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (original) A compound of formula (I)



the *N*-oxides, the pharmaceutically acceptable acid addition salts and the stereochemically isomeric forms thereof, wherein

R¹ is hydrogen, C₁₋₄alkyl, halo, or polyhaloC₁₋₄alkyl;

R² is hydrogen, C₁₋₄alkyl, halo, or polyhaloC₁₋₄alkyl;

R³ is hydrogen or C₁₋₄alkyl;

R⁴ is hydrogen, C₁₋₄alkyl, or halo;

n is an integer 0, or 1;

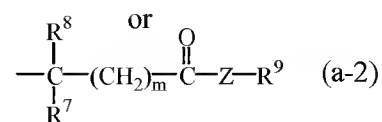
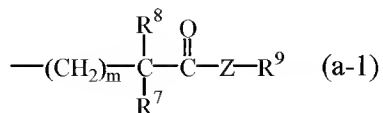
X¹ is carbon and X² is carbon; or X¹ is nitrogen and X² is carbon;

or X¹ is carbon and X² is nitrogen;

X³ is carbon or nitrogen;

Y represents O, or NR⁶ wherein R⁶ is hydrogen or C₁₋₄alkyl;

R⁵ represents a radical of formula



wherein

m is an integer 0, 1, or 2;

Z is O or NH;

R⁷ is hydrogen,

C₁₋₆alkyl;

C₁₋₆alkyl substituted with hydroxy, amino, mono- or di(C₁₋₄alkyl)amino,

C₁₋₄alkyloxycarbonyl, aminocarbonyl, aryl or heteroaryl;

C₁₋₄alkyl-O-C₁₋₄alkyl;

C₁₋₄alkyl-S-C₁₋₄alkyl; or

aryl;

R⁸ is hydrogen or C₁₋₆alkyl;

R⁹ is hydrogen, C₁₋₄alkyl, aryl¹, or C₁₋₄alkyl substituted with aryl¹;

or when Y represents NR⁶ the radicals R⁵ and R⁶ may be taken together with the nitrogen to which they are attached to form pyrrolidinyl substituted with C₁₋₄alkyloxycarbonyl and optionally further substituted with hydroxy; or piperidinyl substituted with C₁₋₄alkyloxycarbonyl;

aryl is phenyl; phenyl substituted with one, two or three substituents each independently selected from C₁₋₄alkyl, C₁₋₄alkyloxy, halo, hydroxy, nitro, cyano, C₁₋₄alkyloxycarbonyl, trifluoromethyl, or trifluoromethoxy; or benzo[1,3]dioxolyl;

aryl¹ is phenyl; phenyl substituted with one, two or three substituents each independently selected from C₁₋₄alkyl, C₁₋₄alkyloxy, halo, hydroxy, nitro, cyano, C₁₋₄alkyloxycarbonyl, trifluoromethyl, or trifluoromethoxy; and heteroaryl is imidazolyl, thiazolyl, indolyl, or pyridinyl.

2. (original) A compound as claimed in claim 1 wherein X¹, X² and X³ are carbon.
3. (original) A compound as claimed in claim 1 wherein R¹ is trifluoromethyl; R² is hydrogen; R³ is hydrogen; R⁴ is hydrogen; X¹, X² and X³ are carbon; n is the integer 1; Y represents NR⁶ wherein R⁶ is hydrogen or methyl; and R⁵ is a radical of formula (a-1) wherein m is the integer 0.

4. (original) A compound as claimed in claim 1 wherein R¹ is trifluoromethyl; R² is hydrogen; R³ is hydrogen; R⁴ is hydrogen; X¹, X² and X³ are carbon; n is the integer 1; Y represents NR⁶ wherein R⁶ is hydrogen or methyl; and R⁵ is a radical of formula (a-1) wherein m is the integer 1.
5. (original) A compound as claimed in claim 1 wherein R¹ is trifluoromethyl; R² is hydrogen; R³ is hydrogen; R⁴ is hydrogen; X¹, X² and X³ are carbon; n is the integer 1; Y represents NR⁶ wherein R⁶ is hydrogen or methyl; and R⁵ is a radical of formula (a-2) wherein m is the integer 1.
6. (previously presented) A compound as claimed in claim 1 wherein R¹ is trifluoromethyl; R² is hydrogen; R³ is hydrogen; R⁴ is hydrogen; X¹, X² and X³ are carbon; n is the integer 1; Y represents NR⁶ and R⁵ and R⁶ are taken together with the nitrogen to which they are attached to form pyrrolidinyl substituted with C₁₋₄alkyloxycarbonyl and optionally further substituted with hydroxy, or piperidinyl substituted with C₁₋₄alkyloxy-carbonyl.
7. (previously presented) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a therapeutically active amount of a compound as claimed in claim 1.
8. (previously presented) A process for preparing a pharmaceutical composition as claimed in claim 7 wherein a therapeutically active amount of a compound as claimed in claim 1 is intimately mixed with a pharmaceutically acceptable carrier.
- 9 – 15. (Cancelled)